Appendix F

Situations for which expert consultation for HIV postexposure prophylaxis is advised

- Delayed (i.e., later than 24-36 hrs.) exposure report
 - o o the interval after which there is no benefit from post exposure prophylaxis (PEP) is undefined
- Unknown source (i.e., needle in sharps disposal container or laundry)
 - o o decide use of PEP on a case by case basis
 - o o consider the severity of the exposure and the epidemiologic likelihood of HIV exposure
 - o o do not test needles or other sharp instruments for HIV
- Known or suspected pregnancy in the exposed person
 - o o does not preclude the use of optimal PEP regimes
 - o o do not deny PEP solely on the basis of pregnancy
- Resistance of the source virus to antiretroviral agents
 - o influence of drug resistance on transmission risk is unknown
 - selection of drugs to which the source person's virus is unlikely to be resistant is recommended, if the source person's virus is known or suspected to be resistant to ≥1 of the drugs considered for the PEP regimen
 - o resistance testing of the source person's virus at the time of the exposure is not recommended
- Toxicity of the initial PEP regimen
 - o o adverse symptoms, such as nausea and diarrhea are common with PEP
 - o symptoms often can be managed without changing the PEP regimen by prescribing antimotility and /or antiemetic agents
 - o modification of dose intervals (i.e., administering a lower dose of drug more frequently throughout the day, as recommended by the manufacturer), in other situations might help alleviate symptoms